

closely related to the target organ damage. It can be provided the basis for the intervention of cardiovascular risk.

<http://dx.doi.org/10.1016/j.hkijn.2015.09.147>

0051

Relationship Between Serum Level of Fibroblast Growth Factor-23 (FGF-23) and Calcium and Phosphate Metabolism of Patients with Chronic Kidney Disease

Lyu Li, Wang Caili, Mi Yan

The First Affiliated Hospital of Baotou Medical College, Baotou, China

Objective: To investigate the relationship between serum level of fibroblast growth factor-23 (FGF-23) and calcium and phosphate of patients with chronic kidney disease.

Methods: Intact FGF-23 and 1,25(OH)₂ vitamin D₃ were detected by enzyme-linked immunosorbent assay (ELISA) in CKD 151 patients with various degree of renal function and fourteen healthy controls, at the same time, some other parameters including creatinine, serum phosphate, calcium, parathyroid hormone, uric acid and alkaline phosphatase were measured. Further, the relationship between FGF-23 and other biochemical variables was analysed.

Results: (1) The level of serum FGF-23 was gradually elevated at CKD stage 1, 2–3, 4–5, there were significant difference in CKD stage 2–3 and 4–5 compared respectively with normal controls ($P < 0.01$), and in CKD stage 4–5 compared with CKD stage 1 and 2–3 ($P < 0.01$). (2) In the CKD stage 1–5 the level of serum FGF-23 by Pearson relativity analysis was significantly correlated positively with creatinine ($r = 0.97$, $P < 0.01$), phosphate ($r = 0.65$, $P < 0.01$), and parathyroid hormone ($r = 0.536$, $P < 0.01$). The level of serum FGF-23 by Pearson relativity analysis was significantly correlated negatively with eGFR ($r = -0.578$, $P < 0.01$), 1,25(OH)₂ vitamin D₃ ($r = -0.586$, $P < 0.01$). (3) In the CKD stage 1–5, creatinine, 1,25(OH)₂ vitamin D₃ and parathyroid hormone were significant variables that influenced the level of serum FGF-23 in multiple regression analysis.

Conclusion: FGF-23 concentrations were gradually elevated in three groups of patients with CKD, serum phosphorus, 1,25(OH)₂D₃, intact PTH and creatinine are the major regulators that influenced FGF-23.

0051 Table 2 151例CKD患者FGF-23与相关变量的相关分析			
N = 151	r-value	P-value	
CREA	0.971	< 0.01	
eGFR	-0.578	< 0.01	
1.25(OH) ₂ D ₃	-0.586	< 0.01	
Ca	-0.327	< 0.01	
P	0.650	< 0.01	
Ca*P	0.477	< 0.01	
UA	0.173	0.033	
ALP	0.143	0.079	
PTH	0.536	< 0.01	

<http://dx.doi.org/10.1016/j.hkijn.2015.09.148>

0051

Table 1 CKD1-5期患者血清各指标的比较($\bar{x} \pm s$)

	CREA ($\mu\text{mol/L}$)	eGFR (ml/min/1.73 m^2)	FGF-23 (pg/ml)	1.25(OH) ₂ D ₃ (pg/ml)	Ca (mmol/L)	P (mmol/L)	Ca*P (mmol^2/L^2)	PTH (pg/L)
正常对照组 N = 14 ($\bar{x} \pm s$)	43.8 \pm 2.71	118.81 \pm 5.42	20.82 \pm 1.20	595.11 \pm 10.03	2.27 \pm 0.19	1.15 \pm 0.13	2.61 \pm 0.43	33.81 \pm 1.47
CKD1期 N = 22 ($\bar{x} \pm s$)	69.82 \pm 13.20	112.62 \pm 8.18	71.19 \pm 8.26	546.78 \pm 102.66	2.27 \pm 0.17	1.23 \pm 0.15	2.79 \pm 0.41	38.59 \pm 3.29
CKD2-3期 N = 47 ($\bar{x} \pm s$)	139.90 \pm 35.19	58.97 \pm 16.96*	81.73 \pm 13.23*	531.12 \pm 114.74	2.24 \pm 0.22	1.35 \pm 0.15*	3.02 \pm 0.40	68.14 \pm 30.13
CKD4-5期 N = 82 ($\bar{x} \pm s$)	440.74 \pm 331.36* Δ	15.75 \pm 7.67* Δ	198.05 \pm 111.68* Δ	409.03 \pm 130.51* Δ	2.10 \pm 0.21	1.81 \pm 0.13* Δ	3.80 \pm 0.43	225.41 \pm 172.42* Δ
方差分析	F 28.67	593.98	38.46	19.9	7.22	209.01	26.44	69.22
	P < 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01

*和对照组比较, $P < 0.05$; Δ 与CKD1、CKD2-3比较, $P < 0.05$.

0052

Function of Reduced Blood Pressure by Compound a-Ketoacid in Non-dialysis Patients with CKD Stage 3–5

Li SongYang

Jinan Central Hospital, Shanghai, China

Objective: Evaluate the effect of blood pressure-lowering by a-ketoacid in patients with CKD.

Methods: Selected non-dialysis patients who had hypertension of CKD 3–5 grade, except three cases. All of the patients had taken anti-hypertension medicine over 3 months. They were treated for an average of 34.5 ± 38.7 months, but SBP was still ≥ 140 mmHg and DBP ≥ 90 mmHg. A total of 102 patients were included in the statistical analysis. Among them, 61 were male and 41 were female. Mean age was 70.78 ± 12.34 years (range, 27 to 86 years). 51 cases were CKD stage 3, 31 cases were CKD stage 4, 20 cases were CKD stage 5. Mean duration of chronic renal insufficiency was 32.7 ± 33.3 months (range, 4 to 120 months). Mean duration of hypertension was 232.4 ± 174.4 months (range, 4 to 636 months). A low protein diet (0.6 g/kg/d) and compound a-ketoacid (2.52 g, tid) were given to all patients. At the same time, the anti-hypertension drugs were not changed during the study. After 6 months, statistical analysis was used to compare blood pressure, PTH and renal function changes before and after treatment.

Results: In 102 cases, total efficiency rate was 83.3%. Mean systolic blood pressure was 163.4 ± 15.8 mmHg before treatment and 137.8 ± 14.8 mmHg after treatment. Mean diastolic blood pressure was 87.9 ± 12.9 mmHg before treatment and 77.0 ± 9.6 mmHg after treatment ($P < 0.01$). Mean systolic blood pressure decreased 25.9 ± 14.8 mmHg and diastolic blood pressure decreased 10.8 ± 9.7 mmHg. There were statistically significant differences. PTH had markedly decreased after treatment, but there was no statistically significant differences in renal function before and after treatment.

Conclusion: Low protein diet combining compound a-ketoacid may help to control hypertension in CKD patients.

<http://dx.doi.org/10.1016/j.hkijn.2015.09.149>

0055

Risk of Death and Cardiovascular Outcomes in Chronic Kidney Disease Patients with Chronic Obstructive Pulmonary Disease

Ping-Hsun Wu¹, Yi-Ting Lin², Ming-Yen Lin^{1,3}, Hui-Min Hsieh⁴, Feng-Shiuan Jian⁵, Mei-Chuan Kuo^{1,3}, Shang-Jhy Hwang^{1,3}, Hung-Chun Chen^{1,3}

¹Division of Nephrology, Department of Internal Medicine, Kaohsiung

Medical University Hospital, Kaohsiung, Taiwan

²Department of Family Medicine, Kaohsiung Municipal Hsiao-Kang Hospital, Kaohsiung, Taiwan

³Faculty of Renal Care, College of Medicine, Kaohsiung Medical University, Kaohsiung, Taiwan

⁴Department of Public Health, Kaohsiung Medical University, Kaohsiung, Taiwan

⁵Institute of Public Health, National Yang-Ming University, Taipei, Taiwan

Objective: Chronic obstructive pulmonary disease (COPD) increases all-cause mortality and cardiovascular (CV) events in general population. However, rare study investigates the death and CV risks among chronic kidney

disease (CKD) with COPD. This population-based cohort study aimed to evaluate the all-cause mortality and major CV events of COPD among CKD patients using national health insurance (NHI) database in Taiwan.

Methods: CKD patients enrolled in the pre-end stage renal disease (pre-ESRD) care management program, patients ≥ 40 years old who had at least one outpatient diagnosis code (ICD-9 codes 585, 581.9) or at least one inpatient diagnosis code and free of any dialysis or renal transplant from 1 January 2007 to 31 December 2011 were included. CKD patients with COPD were matched to control subjects without COPD at a 1:1 ratio using propensity scores. Cox proportional hazards regression models were used to estimate hazard ratios (HRs) for all-cause mortality and major CV events analysis.

Results: Propensity score matching yielded a cohort balanced on age, sex, urbanization, socioeconomic status, enrolled year, comorbidities, Charlson Comorbidity Index score, clinical laboratory data, and medications used, with 2498 CKD patients in the COPD group and 2498 in the non-COPD group. CKD patients with COPD were associated with a higher risk for death (HR, 1.134; 95% CI, 1.037–1.241) and non-significant risk for major CV events (HR, 1.041; 95% CI, 0.905–1.198) compared to those without COPD.

Conclusion: CKD patients with COPD are associated with higher all-cause mortality and might increase risk of major CV events.

<http://dx.doi.org/10.1016/j.hkjin.2015.09.150>

0086

Association Between Anemia and Abnormal Bleeding Times in Patients with Chronic Kidney Disease

Ha Yeon Kim

Chonnam National University Medical School, Gwangju, Republic of Korea

Objective: Platelet dysfunction associated hemorrhagic complications and chronic anemia are often encountered in patients with chronic kidney disease. The present study aimed to evaluate the association of anemia and abnormal bleeding time in patients with chronic kidney disease.

Methods: We retrospectively analyzed the data of 1022 (60.24 ± 15.89 years, men 58.7%), patients with chronic kidney disease.

Results: Patients with prolonged bleeding time (bleeding time > 182 sec) were older age and lower eGFR, Hgb, Hct and PLT, and higher BUN and Cr levels compared to those with normal bleeding time. The prevalence of abnormal bleeding time was found to be higher as Hgb declined [25.0% for patients with Hgb ≥ 10 mg/dl ($n = 595$), 31.9% for patients with $8 \leq$ Hgb < 10 mg/dl ($n = 343$), and 43.4% for patients with Hgb < 8 mg/dl ($n = 84$), $P < 0.001$, respectively]. eGFR was decreased as Hgb declined (29.8 ± 19.0 mL/min/1.73 m² in patients with Hgb ≥ 10 mg/dl, 16.0 ± 13.0 mL/min/1.73 m² in patients with $8 \leq$ Hgb < 10 mg/dl, and 10.3 ± 8.61 mL/min/1.73 m² in patients with Hgb < 8 mg/dl, $P < 0.001$, respectively). Multivariate analysis revealed that age (OR, 1.019; 95% CI, 1.010–1.029), $8 \text{ mg/dl} \leq \text{Hgb} < 10 \text{ mg/dl}$ (OR, 1.214; 95% CI, 0.873–1.686), Hg $< 8 \text{ mg/dl}$ (OR, 1.820; 95% CI, 1.084–3.055), $15 \text{ mL/min/1.73 m}^2 \leq \text{eGFR} < 30 \text{ mL/min/1.73 m}^2$ (OR, 1.616; 95% CI, 1.053–2.481), eGFR $< 15 \text{ mL/min/1.73 m}^2$ (OR, 1.668; 95% CI, 1.160–2.397), thrombocytopenia (PLT $< 150 \times 10^9/\text{L}$) (OR, 2.659; 95% CI, 1.981–3.569) were independently associated with prolonged bleeding time.

Conclusion: Severe anemia (Hgb < 8 mg/dl) is independently associated with prolonged bleeding time in chronic kidney disease, even after adjusting for eGFR and other potential confounders.

<http://dx.doi.org/10.1016/j.hkjin.2015.09.151>

0087

Low Plasma Level of Cathelicidin is Associated with Decreased eGFR

Ha Yeon Kim, Seung Jin Lee

Chonnam National University Medical School, Gwangju, Republic of Korea

Objective: This study assessed plasma cathelicidin, 25-OH vitamin D, and natural killer (NK) cells which play an important role in innate immunity.

Methods: The study cohort included 175 patients who have a variety of estimated glomerular filtration rate (eGFR) at the Chonnam National University Hospital. Plasma cathelicidin level was measured by ELISA and absolute

count of NK cell was determined by flow cytometry. NK cell was identified phenotypically as CD3-CD56+.

Results: The study population was divided into 3 groups according to the eGFR: Group I, eGFR ≥ 60 mL/min/1.73 m², $n = 33$ (18.9%); group II, $15 \leq$ eGFR < 60 mL/min/1.73 m², $n = 47$ (26.8%); group III, eGFR < 15 mL/min/1.73 m², $n = 95$ (54.3%), respectively. Plasma cathelicidin level was decreased with lower eGFR (189.3 ± 18.21 ng/ml in group I, 182.6 ± 17.69 ng/ml in group II, and 179.7 ± 17.76 ng/ml in group III, $P = 0.032$, respectively). On the other hand, percentages and absolute numbers of NK cells were significantly higher in the peripheral blood of patients with lower eGFR (1897.7 ± 1584.05 in group I, 2110.8 ± 2168.37 in group II, and 2833.8 ± 2149.65 in group III, $P = 0.034$, respectively). A 25-OH vitamin D was not distinguished in these groups, but after an exclusion of patients with hemodialysis, 25-OH vitamin D level showed decreasing tendency with lower eGFR (16.4 ± 6.22 ng/ml in group I, 12.4 ± 4.74 ng/ml in group II, and 11.2 ± 4.96 ng/ml in group III, $P = 0.166$, respectively).

Conclusion: Plasma cathelicidin level was significantly decreased with lower eGFR while an absolute count of NK cells was increased with lower eGFR, which might be associated with dysfunction of NK cells. After an exclusion of patients with hemodialysis, 25-OH vitamin D concentration showed decreasing tendency with lower eGFR.

<http://dx.doi.org/10.1016/j.hkjin.2015.09.152>

0088

Calciophylaxis and Hydroxyapatite Crystal Deposition in a Patient with Uremia

Wu Yongyao

Beilun Branch of the First Affiliated Hospital, Zhejiang University School of Medicine, Ningbo, China

A 42-year-old male patient with uremia on hemodialysis was admitted due to right foot ulcer and infection. Two years ago, he lost sight in the right eye due to calcium salt deposition. One year ago, he developed multiple peri-articular masses with progressive enlargement, which were identified as hydroxyapatite. Acral gangrene occurred repeatedly throughout the course of disease. Radiology indicated diffuse and continuous calcification of middle and small arteries. Concomitant calcific uremic arteriopathy and hydroxyapatite crystal deposition disease were considered. The diseases originated from abnormal disposition of hydroxyapatite in conjunctiva, vessels and articular peripheries and shared a common pathogenesis.

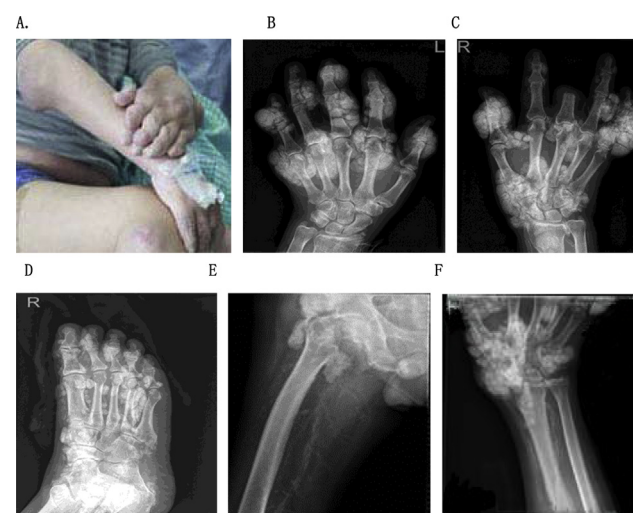


Fig.1 A. Multiple juxta-articular masses near bilateral elbow joints, wrist joints, metacarpophalangeal joints, interphalangeal joints, toe joints and knee joints. B, C, D, E, F. X-ray manifestations: mass-like calcification shadows in the peripheries of multiple joints all over the body; subcapital fracture of right femur; linear calcification of middle and small arteries of limbs, calcification of reticular vessels.